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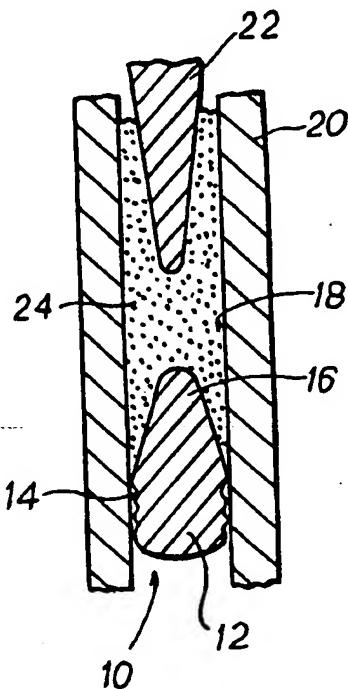
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(54) Plugs for the medullary canal of a bone

(57) An implant 22, e.g. for locating a hip prosthesis, is placed in the medullary canal 18 of a bone 20 (e.g. the femur) and secured by pressurised cement 24. A push-fit plug 10 prevents cement penetrating down the canal. The plug can be biodegradable so that in time it dissolves away and does not modify the flexural rigidity of the cement/bone system and the risk of fractures occurring at the level of the plug is reduced. Additionally or alternatively, the outwardly facing end portion 16 of the plug may be generally conical so that the thickness of the cement increases gradually and the flexural rigidity likewise changes gradually.

FIG. 1



20 18 16 12 10

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FIG. 1

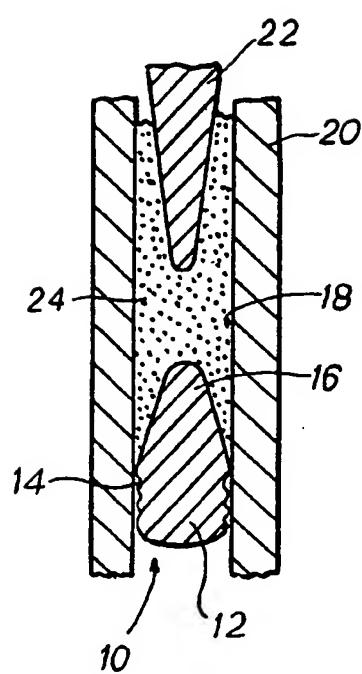
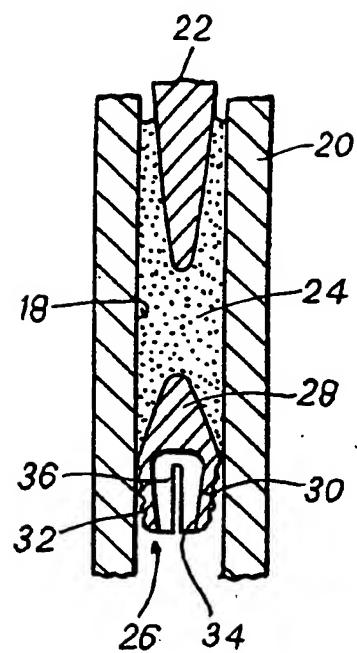


FIG. 2



SPECIFICATION**Plugs for the medullary canal of a bone**

5 This invention relates to surgery in which an implant is placed in the medullary canal of a bone, and more particularly to plugs for insertion in the medullary canal in such surgery.

An implant has to be placed in the medullary canal 10 of a femur when a patient is to be given an artificial hip joint, in order to locate the hip prosthesis. The implant is located in the medullary canal by use of a bone cement, e.g. polymethylmethacrylate cement. Such cement does not have any appreciable adhesive qualities, and relies for its security on obtaining a mechanical interlock with irregularities in the bone. Such mechanical interlocking is greatly enhanced if the cement is introduced into the canal under pressure. However, there is a tendency for the cement to 15 flow down the medullary canal so it is not possible to use much pressure. To overcome this, it is known to insert a plug which is a push fit inside the medullary canal and which prevents the cement flowing down the canal and permits greater pressurisation of the 20 cement. It is known for such plugs to be formed from bone, or from bone cement. An improved plug made from a plastics material is the Seidel plug. Such a plastics plug will of course remain inside the medullary canal when the surgery is completed. A problem 25 with all such known plugs is that because of such factors as the difference in Young's modulus between the bone and the bone cement, there is an abrupt change in the flexural rigidity of the bone at the plug. This has not uncommonly led to fractures 30 at the level of the plug, for example if the patient should suffer a fall. A fracture at this level is very difficult to set.

According to one aspect of the present invention we provide a plug for the medullary canal of a bone, 40 having a portion with sides adapted to be a push fit in the medullary canal, the plug being made of a biodegradable material.

Preferably the plug has a tapering end portion, which preferably is generally conical. The plug can 45 then be inserted in the medullary canal with the conically tapering portion facing into the part of the canal into which cement is to be introduced, so that when the cement is introduced the conically tapering portion forms a mould so that there is a gradually 50 increasing thickness of cement in contact with the bone along the length of the tapering portion. This can then provide a gradual change in flexural rigidity of the cement/bone system along the plug. Being biodegradable, the plug will eventually dissolve, and 55 so have no effect on flexural rigidity.

According to a second aspect of the invention we provide a plug for the medullary canal of a bone, comprising a portion having sides adapted to be a push fit in the medullary canal, and a tapering end 60 portion, preferably a generally conically tapering end portion. This plug can be of plastics material. The portion which has sides adapted to be a push fit in the medullary canal may be hollow, and may also have one or more slits in a longitudinal plane which 65 permit flexing of said sides to facilitate insertion in

the canal.

The invention will be more clearly understood by reference to the accompanying drawings, wherein Figs. 1 and 2 are schematic sectional views of two

70 plugs embodying the invention, *in situ* in a bone. Referring to Fig. 1, the plug 10 has a generally cylindrical portion 12 with generally parallel serrated sides 14. The sides 14 merge into a generally conically tapered portion 16 of the plug. During surgery, 75 the medullary canal 18 of the bone 20 such as a femur will be reamed out, and a plug 10 of suitable size for the reamed out canal will be introduced with the cylindrical portion 12 leading so that the conically tapering portion 16 faces outwardly. The portion 80 12 is a push fit inside the canal 18, assisted by the serrations 14, and the plug is inserted for a sufficient distance to permit filling of the canal with cement as desired.

Bone cement 24, e.g. polymethylmethacrylate 85 bone cement, is introduced into the open end of the medullary canal 18, suitably by retrograde filling above the plug 10, and pressurised. The desired implant 22 such as a hip prosthesis is then introduced into the medullary canal 18 by being inserted 90 down the centre of the previously placed bone cement. Further pressurisation of the bone cement is achieved at this stage by occluding the top of the medullary canal and allowing the generally wedge-shaped stem of the implant to force bone cement 95 further into the medullary canal of the femur. Pressurisation of the cement ensures that it is forced into the inter trabecular spaces of the bone structure thereby providing a good mechanical interlock so as to anchor the implant securely. The plug 10 prevents pressurised cement flowing further down the medullary canal.

Because the conically tapering portion 16 of the plug faces into the reamed out cavity of the canal 18 which is filled with cement, it forms a mould for the 105 cement so that the thickness of cement which is in contact with and mechanically interlocking with the bone 20 increases gradually along the length of the tapering portion 16. This means that the flexural rigidity of the cement/bone system also changes 110 gradually in this region. The risk of the bone fracturing in this region is therefore reduced.

The plug 10 is manufactured from a biodegradable material comprising stabilised ox fibrin, mixed with 35% glycerol as a plasticizer. This material is 115 described by Ian Cappernauld, P. Lawrie and D. A. French in "Properties of Bovine Fibrin Absorbable Implants", *Surgery: Gynecology & Obstetrics*, January 1977, Vol. 144, 3-7. It is available from Ethnor Division, Ethicon Limited, P.O. Box 408, Bankhead 120 Avenue, Edinburgh, EH11 4HE, under the Trade Mark BIETHIUM. The plug 10 can be made by compression moulding the powdered material, or by machining from a round bar. Other biodegradable materials could of course be used. Because it is made of 125 biodegradable material, the plug 10 eventually dissolves, leaving behind the cement 24 with its gradually increasing thickness around the region once occupied by the portion 16. Once this has happened, the flexural rigidity of the cement/bone system is 130 unmodified by the plug, and so the risk of a fracture

occurring in this area is reduced even further.

- Referring now to Fig. 2, a plug 26 is shown which is moulded in the usual way from a non-biodegradable plastics material. A suitable material 5 is ultra-high molecular weight polyethylene. It is generally similar to the plug 10 shown in Fig. 1, and in particular has a conically tapering portion 28 which behaves in exactly the same manner as in Fig. 1. There is also a generally cylindrical portion 30 10 which has serrated sides 32 which are a push fit in the medullary canal 18, as before. However, the portion 30 is hollowed out to give a recess 34 open at the end of the plug remote from the portion 28. Two slits 36 are provided through the walls of the portion 15 30, the slits being open at the end of the plug remote from the portion 28, and being in a longitudinal plane with respect to the plug. This effectively divides the portion 30 into two halves which can flex slightly towards each other. This facilitates insertion 20 of the plug 26 into the canal 18.

CLAIMS

1. A plug for the medullary canal of a bone, comprising a first portion having sides adapted to be a push fit in the medullary canal, and a second portion 25 at one end thereof which tapers in the direction away from the first portion.
2. A plug according to claim 1 made from plastics material.
3. A plug according to claim 1 made from a 30 biodegradable material.
4. A plug according to claim 3 wherein the biodegradable material comprises ox fibrin.
5. A plug according to claim 4 wherein the 35 biodegradable material contains glycerol as a plasticizer.
6. A plug according to claim 1 wherein the first portion is hollow.
7. A plug according to claim 6 wherein the hollow first portion has one or more slits in a longitudinal plane which permit flexing of said sides to facilitate insertion in the canal.
8. A plug according to claim 1 wherein the tapering second portion is generally conical.
9. A plug for the medullary canal of a bone, having 45 a first portion with sides adapted to be a push fit in the medullary canal, wherein the plug is made of a biodegradable material.
10. A plug according to claim 9 wherein the biodegradable material comprises ox fibrin.
- 50 11. A plug according to claim 10 wherein the biodegradable material contains glycerol as a plasticizer.
12. A plug for the medullary canal of a bone, substantially as described herein with reference to 55 Fig. 1 or Fig. 2 of the accompanying drawings.